Parathyroid Hormone Analogs
Evenity (romosozumab-aggg), Forteo (teriparatide), Tymlos (abaloparatide)

Evenity (romosozumab-aqqg), i orteo (temparatide), i yimos (abaioparatide)					
Member and Medication Information (required)					
Member ID:			Member Name:		
DOB:			Weight:		
Medication Name/ Strength:			Dose:		
Dire	ections for use:				
		Provider Infor	mation (required		
		NPI:		Specialty:	
Contact Person:		Office Phone:		Office Fax:	
	FAX FORM AND RELEVA CHART NOTES and/or U			ABORATORY RESULTS, SSITY TO 855-828-4992	
Crite	ria for Approval (all of the following m	ust be met):			
	Diagnosis of one of the following:				
[Postmenopausal osteoporosis. Char	teoporosis. Chart note page #:			
[Osteoporosis due to sustained systems	·	erapy. (Forteo only). C	hart note page #:	
Į	•	Osteoporosis due to primary hypogonadism in males. (Forteo only). Chart note page #:			
u ,	Very High risk for fracture defined as:	ery High risk for fracture defined as:			
Į	Intolerance to antiresorptive therapy (bisphosphonates, denosumab) OR				
 Osteoporotic fracture while on antiresorptive therapy OR Previous osteoporotic fracture OR ≥ 40 years old with one of the following: T-score ≤ -2.5 at the femoral neck or spine T-score between -1.5 and -2.0 with a 10-year probability of major osteoporotic fracture ≥ 20% or hip fracture ≥ 3% 					
				ic fracture ≥ 20% or hip fracture ≥ 3%	
Note					
 Boxed warning: Parathyroid hormone analogs are not recommended for use in patients with increased risk for osteosarco 					
	Goved warning: Parathyroid normone analogs are not recommended for use in patients with increased risk for osteosarcoma (e.g. Paget's disease of bone, unexplained elevations of alkaline phosphatase, open epiphyses, bone metastases or skeletal malignancies, hereditary disorders predisposing to osteosarcoma, or prior external beam or implant radiation therapy involving the skeleton) or who have an underlying hypercalcemic disorder (e.g. primary hyperparathyroidism).				
* 1	Boxed warning: Evenity (romosozumab-aqqg) may increase the risk of myocardial infarction, stroke, and cardiovascular death.				
It should not be initiated in patients who have had a myocardial infarction or stroke v				- ·	
				ctors. If a patient experiences a myocardial	
İ	infarction or stroke during therapy, ther	apy should be discont	inued.		
Initia	al Authorization: Up to 12 months for Ev	venity, up to 24 month	s for Forteo and Tymlo	os.	
	uthorization: None, lifetime limits apply		,		
PROVIDER CERTIFICATION					
I her	eby certify this treatment is indicated, n	ecessary and meets th	e guidelines for use.		
Dross	criber's Signature		 Date		
ries(LINEI 9 SIKIIALUIE		Date		